

AD No. 21 / 47

ASTIA FILE COPY

Report No. 1

on

CONTRACT NO. Nmr-495(04)  
NF 055 398

Shalom, David, and Melvin S. Newman  
October 1, 1953

THE OHIO STATE UNIVERSITY  
RESEARCH FOUNDATION

BEST AVAILABLE COPY

RF Project 497

Report No. 1

TECHNICAL

R E P O R T

by

THE OHIO STATE UNIVERSITY  
RESEARCH FOUNDATION

Columbus 10, Ohio

To

OFFICE OF NAVAL RESEARCH  
Contract No. Nonr-495(04)  
NR 055 298

On

STERIC FACTORS IN ORGANIC CHEMISTRY

Submitted by

Shalom Sarel and Melvin S. Newman  
Department of Chemistry

Date October 1, 1953

## SUMMARY

The synthesis of a number of acids, alcohols derived by reduction, and the acetates of these alcohols is described. Physical properties and yields are tabulated.

The rates of alkaline hydrolysis of these acetates at 20° and 30° in aqueous dioxane have been measured and calculations of rate constants and thermodynamic properties have been made.

# ALKALINE HYDROLYSIS OF ACETATES OF HIGHLY BRANCHED ALCOHOLS

## INTRODUCTION

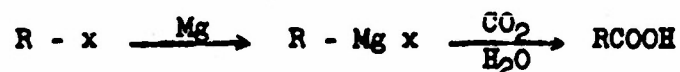
The over-all objective of the work herein reported is to accumulate quantitative data on the saponification of a large number of primary, secondary, and tertiary acetates. These data will be used in a study of steric effects of various alkyl groups in alkaline hydrolysis. It is hoped that eventually these data will be useful in predicting the alkyl ester to be used in many different reactions where bifunctional compounds, one function of which is the ester function, are involved.

## EXPERIMENTAL

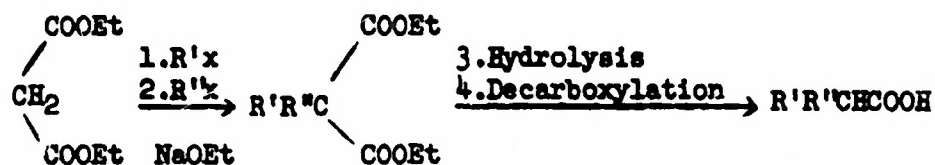
### A. SYNTHESIS OF ACIDS

Five different methods were used for the preparation of the carboxylic acids or their esters. These methods are outlined below.

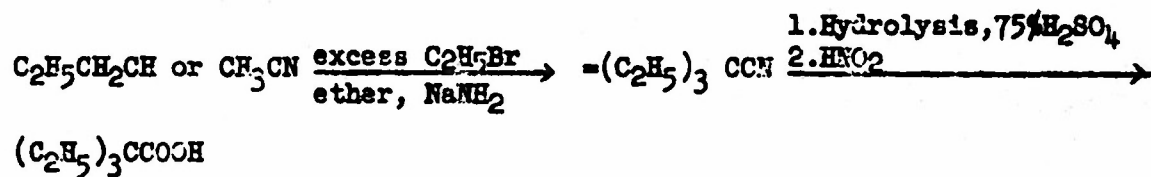
#### 1. Carboxylation of the Grignard reagent:



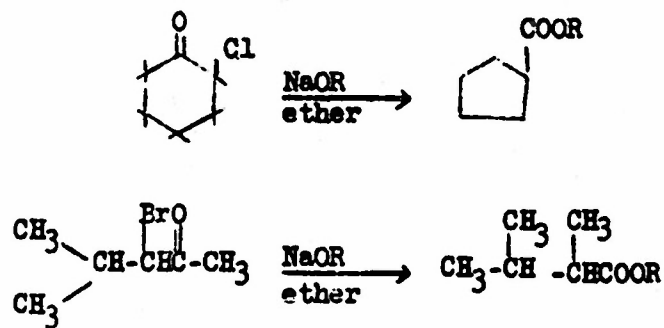
2. Alkylation of Malonic or Cyanoacetic Esters:



3. Alkylation of Nitriles (Ziegler Method):



4. Alkaline Rearrangement of  $\alpha$ -Haloketones (Favorski Reaction):



5.



## B. REDUCTION OF ACIDS AND ESTERS

The use of  $\text{LiAlH}_4$  as a reducing reagent for converting carboxylic acids and esters into the corresponding primary alcohols has been found highly satisfactory.

## C. FORMATION OF ACETATES

The primary alcohols were easily acetylated either by (1) acetyl chloride or (2) acetic anhydride in presence of dry pyridine, affording very good yields, depending on effectiveness of isolation methods of the product used. The data are summarized in Table I. The physical properties are listed in Table II.

The secondary and tertiary alcohols were acetylated by acetyl chloride and magnesium metal in dry ether as follows: (See Table III).



## D. RATES OF SAPONIFICATION

The rate of hydrolysis of the acetates with 0.01 N sodium hydroxide, using water or 70% aqueous dioxane as solvents, was followed titrimetrically at 20°C and 30°C. The rate constants are given in Table IV. The thermodynamic data in Table IV were calculated using the equations listed below.

#### E. CALCULATION OF ARRHENIUS PARAMETERS

$$(1) \Delta E^* = 2.303 \times R \times \frac{T_1 \cdot T_2}{T_2 - T_1} \cdot \log \frac{k_2}{k_1} \quad (\text{Energy of activation})$$

$$(2) \log PZ = \log k + \frac{\Delta E^*}{2.303 \cdot RT} = B$$

$$(3) \Delta S^* = 2.303 \times R \left( B - \log \frac{RT}{Nh} \right) \quad (\text{entropy of activation for all degrees of freedom})$$

which are derived from the transition state theory of reaction velocity:

$$k = \frac{RT}{Nh} \cdot e^{-\Delta S^*/R} \cdot e^{-\Delta E^*/RT}$$

$$\text{or} \quad \log k = \log \frac{RT}{Nh} + \frac{\Delta S^*}{2.303 R} - \frac{\Delta E^*}{2.303 RT}$$

where  $R/N$  and  $h$  are the Boltzman and Planck constants, respectively, and  $\Delta S^*$  is the entropy of activation for all degrees of freedom.

Equation (2) is used for calculating  $B (= \log PZ)$  at each temperature; average  $B$  is used in equation (3). In equation (3) the average temperature is used as  $T = 298$ .

#### F. INFRARED SPECTRA

Spectra were recorded between 5000 and 625  $\text{cm}^{-1}$  with a Baird Infrared Recording Spectrophotometer, Model B. All liquids were

measured in a sandwich-type sodium chloride cell. See data in Table II.

#### DISCUSSION OF RESULTS

As stated before, one of the prime purposes of this study is to obtain a quantitative measure of steric hindrance towards the alkaline hydrolysis of alkyl acetates. In order to rule out polar effects as much as possible, comparisons of relative rates will be made only within the members of the same class of esters. That is, the rates of primary alkyl esters will form one class, the rates of secondary esters another, and the rates of tertiary esters still another. It is hoped that a knowledge of the magnitude of the steric factors involved will prove useful in determining that ester of an acid to be used in certain reactions involving bi-functional compounds one function of which is the ester function.

There are certain cases described in the literature in which the use of t-butyl esters is preferable to that of ethyl esters. However, the preparation of t-butyl esters is often quite tedious. It would be helpful to know what primary and secondary alcohols might be expected to show the same degree of hindrance. The completion of our work on the alkaline saponification of primary alkyl acetates allows this to be done, as will be shown below.

The rates for alkaline hydrolysis of methyl, ethyl, isopropyl, and t-butyl acetates are 7.84, 4.57, 1.26, and 0.081 respectively,



using the same units as those indicated in Table IV. Thus we see that compounds 3 and 9 saponify slightly more slowly than t-butylacetate even though primary alkyl groups are involved. All of the other primary alcohol esters listed in Table IV saponify more slowly than methyl and ethyl acetate and hence one has a nice selection of primary alkyl groups having larger steric requirements than ethyl. It remains for much future work to show how valuable this information may prove to be in the selection of the proper ester groups in various cases.

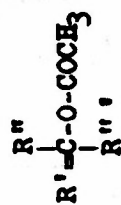
TABLE I. ACETATES OF PRIMARY ALCOHOLS OF FORMULA  $\text{R}^{\text{I}} - \text{C}(\text{R}^{\text{II}})(\text{R}^{\text{III}})\text{COOCH}_3$

No.	Alcohol	R <sup>I</sup>	R <sup>II</sup>	R <sup>III</sup>	R <sup>II</sup>	Acid yield	Ref.	Alcohol yield	Acetate
1.	Isobutyl	CH <sub>3</sub>	CH <sub>3</sub>	H	H	-		-	-
2.	Neopentyl	CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>			91%	80%
3.	2,2-Dimethylbutyl	CH <sub>3</sub>	CH <sub>3</sub>	CH <sub>3</sub>	C <sub>2</sub> H <sub>5</sub>	40%	A-1	90%	92%
4.	2,2-Diethylbutyl	C <sub>2</sub> H <sub>5</sub>	C <sub>2</sub> H <sub>5</sub>	C <sub>2</sub> H <sub>5</sub>	C <sub>2</sub> H <sub>5</sub>	60%	A-3	65%	93%
5.	2-Methylbutyl	CH <sub>3</sub>	C <sub>2</sub> H <sub>5</sub>	H	H				77%
6.	2-Ethylbutyl	C <sub>2</sub> H <sub>5</sub>	C <sub>2</sub> H <sub>5</sub>	H	H				70%
7.	3-Methyl-2-ethylbutyl	C <sub>2</sub> H <sub>5</sub>	(CH <sub>3</sub> ) <sub>2</sub> CH	H	H	78.5%	A-2	83%	90%
8.	3,3-Dimethyl-2-ethylbutyl	C <sub>2</sub> H <sub>5</sub>	(CH <sub>3</sub> ) <sub>3</sub> C	H	H			96%	97%
9.	2-2-Diisopropylethyl	(CH <sub>3</sub> ) <sub>2</sub> CH	(CH <sub>3</sub> ) <sub>2</sub> CH	H	H	50%	A-2	67%	75%
10.	2,3-Dimethylbutyl	(CH <sub>3</sub> ) <sub>2</sub> CH	CH <sub>3</sub>	H	H	60%	A-2 A-4	90%	92%
11.	Cyclopropanemethyl		Cyclopropane	H	H	94.8%	A-5	70%	88.9%
12.	Cyclobutanemethyl		Cyclobutane ring	H	H	-	-	80%	92%
13.	Cyclopentanemethyl		Cyclopentane ring	H	H	65%	A-4	80%	90%
14.	Cyclohexanemethyl		Cyclohexane ring	H	H	-	-	-	77%
15.	2-Ethylhexyl-1	C <sub>2</sub> H <sub>5</sub>	n-C <sub>4</sub> H <sub>9</sub>	H	H	-	-	-	93%

TABLE II. PHYSICAL PROPERTIES AND C-O ABSORPTION BAND IN THE INFRARED  
SPECTRA OF ACETATES OF PRIMARY ALCOHOLS

No.	Compound Acetate of	B.P. °C	Pressure in mm.	n <sub>D</sub> <sup>25</sup>	C-O Absorption	
					cm <sup>-1</sup>	$\mu$
1	2-Methylpropanol	117	741	1.3885	1046	9.56
2	2,2-Dimethylpropanol	127-127.5	741	1.3927	1031 (1050)	9.67 (9.55)
3	2,2-Dimethylbutanol	152-153	740	1.4188	1038	9.64
4	2,2-Diethylbutanol	103-104	40	1.4269	1034 (1055)	9.69 (9.50)
5	2-Methylbutanol	138-139	741	1.4004	1042	9.58
6	2-Ethylbutanol	160	740	1.4090	1038-1041	9.62-9.65
7	3-Methyl-2-ethylbutanol	88-89	30	1.4156	1036	9.66
8	3,3-Dimethyl-2-ethylbutanol	100	40	1.4220	1038	9.65
9	2,3-Dimethylbutanol	147	740	1.4068	1038	9.65
10	Cyclopropanemethanol	133.5	740	1.4156	1034	9.67
11	Cyclobutanemethanol	150	741	1.4245	1033	9.70
12	Cyclopentanemethanol	172.5	741	1.4340	1038	9.65
13	Cyclohexanemethanol	108	40	1.4421	1040	9.63
14	2,2-Dimethylpropylethanol	185-186	740	1.4214	1045	9.62
15	2-Ethylhexanol-1	140	40	1.4182	1041	9.62

TABLE III. ACETATES OF SECONDARY AND TERTIARY ALCOHOLS



No.	Compound	R'	R''	R'''	Car- binol yield	Acce- tate yield	Ref.
16.	Triethyl carbinyl	C <sub>2</sub> H <sub>5</sub>	C <sub>2</sub> H <sub>5</sub>	C <sub>2</sub> H <sub>5</sub>	75%*	70%	Spassow
17.	Di-isopropyl carbinyl	(CH <sub>3</sub> ) <sub>2</sub> CH	H	(CH <sub>3</sub> ) <sub>2</sub> CH		90%	Spassow
18.	tert. Butyl methyl carbinyl	(CH <sub>3</sub> ) <sub>3</sub> C	H	CH <sub>3</sub>	87%**	63%	AcCl+Py

\*Prepared by Grignard method

\*\*Reduction of ketone with LiAlH

Rates of saponification of above acetates in 70% aqueous dioxane were too slow at 20°C, so that the error in the measurement is too great. Further studies on these compounds will be made.

TABLE IV. KINETICS OF ALKALINE HYDROLYSIS  
OF ACETATES OF PRIMARY ALCOHOLS

No.	Compound	$k$ $\frac{\text{g. mole/l.} \cdot \text{min}^{-1}}{20^\circ\text{C}}$	$\Delta E^\ddagger$ Cal./mole	log. PZ	$\Delta S^\ddagger$ Cal./deg./mole
1	$\text{CH}_3-\overset{\text{CH}_3}{\underset{\text{CH}_3}{\text{C}}}-\text{CH}_2\text{OCOCH}_3$	0.280	9925	$7.3079 \pm 0.130$	+ 1.64
2	$\text{C}_2\text{H}_5-\overset{\text{CH}_3}{\underset{\text{CH}_3}{\text{C}}}-\text{CH}_2\text{OCOCH}_3$	0.210	8258	$6.0955 \pm 0.111$	- 3.91
3	$\text{C}_2\text{H}_5-\overset{\text{C}_2\text{H}_5}{\underset{\text{C}_2\text{H}_5}{\text{C}}}-\text{CH}_2\text{OCOCH}_3$	0.050	13280	$9.7773 \pm 0.107$	+12.94
4	$\text{C}_2\text{H}_5-\overset{\text{CH}_2}{\underset{\text{CH}_2}{\text{C}}}-\text{CH}_2\text{OCOCH}_3$	0.470	8670	$6.3766 \pm 0.110$	- 2.62
5	$\text{C}_2\text{H}_5-\overset{\text{C}_2\text{H}_5}{\underset{\text{C}_2\text{H}_5}{\text{C}}}-\text{CH}_2\text{OCOCH}_3$	0.240	12420	$9.1412 \pm 0.163$	+10.03
6	$\text{C}_4\text{H}_9-\overset{\text{C}_2\text{H}_5}{\underset{\text{C}_2\text{H}_5}{\text{C}}}-\text{CH}_2\text{OCOCH}_3$	0.156	14300	$10.4189 \pm 0.311$	+15.88
7	$\text{CH}_3\text{CH}_2-\overset{\text{CH}_3}{\underset{\text{CH}_3}{\text{C}}}-\text{CH}_2\text{OCOCH}_3$	0.350	12400	$9.1063 \pm 0.149$	+ 9.87

Table IV (continued)

No.	Compound	k		$\Delta E^\ddagger$ Cal./mole	$\Delta S^\ddagger$ Cal./deg./mole
		g. mole/l. min <sup>-1</sup> 20°C	30°C		
8	$\text{CH}_3\text{CH}(\text{CH}_3)\text{CH}_2\text{OCOCH}_3$	0.227	0.411	10240	+ 2.73
9	$\text{CH}_3\text{C}(\text{CH}_3)(\text{CH}_2\text{CH}_2\text{OCOCH}_3)\text{CH}_2\text{CH}_3$	0.060	0.109	10700	+ 4.51
10	$\triangle - \text{CH}_2\text{OCOCH}_3$	1.080	2.280	12900	+12.40
11	$\diamond - \text{CH}_2\text{OCOCH}_3$	0.742	1.473	11820	+ 8.29
12	$\square - \text{CH}_2\text{OCOCH}_3$	0.537	1.078	12020	+ 8.66
13	$\square - \text{CH}_2\text{OCOCH}_3$	0.330	0.705	13100	+12.28
14	$\begin{array}{c} \text{CH}_3 \diagup \text{CH} \\   \\ \text{CH}_3 \end{array} \text{CH} - \text{CH}_2\text{OCOCH}_3$	0.120 <sup>x</sup>			

<sup>x</sup>This result is being checked

Signature Page to Technical Report 2  
Contract No. Nonr-495(04) NR 055 298

STERIC FACTORS IN ORGANIC CHEMISTRY

Investigator Shalom Garefman Date 11-9-53

Supervisor Melvin S. Newman Date 11-9-53

FOR THE OHIO STATE UNIVERSITY RESEARCH FOUNDATION

Executive Director Osam C. Woolpert Date 9 Nov. 1953  
W.Z.H.

Distribution List for Technical Reports  
Contract Nonr-495(00) - NR 055-298  
ROLE OF STERIC FACTORS IN ORGANIC CHEMISTRY

31 July 53  
RF 497

Copies

- 1 Director, ONR Branch Office  
150 Causeway Street  
Boston, Mass.
- 2 Director, ONR Branch Office  
The John Crerar Library Bldg.  
10th Floor, 86 E. Randolph St.  
Chicago 1, Illinois
- 1 Director, ONR Branch Office  
346 Broadway  
New York 13, N.Y.
- 1 Director, ONR Branch Office  
1000 Geary Street  
San Francisco 9, Calif.
- 1 Director, ONR Branch Office  
1030 N. Green Street  
Pasadena 1, Calif.
- 2 Officer in Charge  
ONR, Navy No. 100  
Fleet Post Office  
New York, New York
- 6 Director, Naval Research Lab.  
Washington 25, D.C.  
Attn: Technical Information  
Officer
- 2 Chief of Naval Research  
Washington 25, D.C.  
Attn: Chemistry Branch
- 1 Dr. Ralph G.H. Sie, Research  
Director  
General Laboratories, QM Depot  
2800 S. 20th Street  
Philadelphia 45, Pennsylvania
- 1 Dr. Warren Stubblebine  
Research Director  
Chemical and Plastics Section RDB-MPD  
Quartermaster General's Office  
Washington 25, D.C.

Copies

- 1 Dr. A. Stuart Hunter,  
Technical Director  
Research and Development Branch  
Quartermaster General's Office  
Washington 25, D.C. MPD
- 1 Dr. A. Weissler  
Department of The Army  
Office of the Chief of Ordnance  
Washington 25, D.C.  
Attn: ORDTB-PS
- 1 Research and Development Group  
Logistics Division, General Staff  
Department of The Army  
Washington 25, D.C.  
Attn: Dr. W. T. Read  
Scientific Adviser
- 2 Director, Naval Research Lab.  
Washington 25, D.C.  
Attn: Chemistry Division
- 2 Chief of the Bureau of Ships  
Navy Department  
Washington 25, D.C.  
Attn: Code 340
- 2 Chief of Bureau of Aeronautics  
Navy Department  
Washington 25, D.C.  
Attn: Code TD-4
- 2 Chief of the Bureau of Ordnance  
Navy Department  
Washington 25, D.C.  
Attn: Code Rexd
- 1 Dr. H. A. Zahl, Tech. Director  
Signal Corps Engineering Labs.  
Fort Monmouth, New Jersey



31 July 53  
497

Contract Nonr-495(00) - NR 055-298

Copies

- 1 U.S. Naval Radiological  
Defense Lab.  
San Francisco 24, Calif.  
Attn: Technical Library
- 2 Naval Ordnance Test Station  
Inyokern  
CHINA LAKE, California  
Attn: Head, Chemistry Div.
- 1 Office of Ordnance Research  
2127 Myrtle Drive  
Durham, North Carolina
- 1 Technical Command  
Chemical Corps  
Chemical Center, Maryland
- 1 U.S. Atomic Energy Commission  
Research Division  
Washington 25, D.C.
- 1 U.S. Atomic Energy Commission  
Chemistry Division  
Brookhaven National Laboratory  
Upton, New York
- 1 U.S. Atomic Energy Commission  
Library Branch, Tech.Info., ORE  
P.O. Box E  
Oak Ridge, Tennessee
- 1 University of California  
Department of Chemistry  
Los Angeles 24, California  
Attn: Dr. S. Winstein
- 1 Columbia University  
Department of Chemistry  
New York 27, N.Y.  
Attn: Dr. L. P. Hammett
- 1 University of Colorado  
Department of Chemistry  
Boulder, Colorado  
Attn: Dr. S. J. Cristol

Copies

- 1 Harvard University  
Department of Chemistry  
Cambridge 38, Massachusetts  
Attn: Dr. P. D. Bartlett
- 1 Purdue University  
Department of Chemistry  
Lafayette, Indiana  
Attn: Dr. Herbert C. Brown
- 1 Massachusetts Institute of Tech.  
Department of Chemistry  
Cambridge, Massachusetts  
Attn: Dr. C. G. Swain
- 1 Pennsylvania State College  
Department of Chemistry  
State College, Pennsylvania  
Attn: Dr. R. W. Taft, Jr.
- 1 University of Southern California  
Department of Chemistry  
Los Angeles 7, California  
Attn: Dr. R. F. Brown
- 5 ASTIA Document Service Center  
Knott Building  
Dayton 2, Ohio
- 1 Office of Technical Services  
Department of Commerce  
Washington 25, D.C.
- 1 Office of Secretary of Defense  
Pentagon, Room 3D1041  
Washington 25, D.C.  
Attn: Library Branch (R and D)
- 1 Dr. A. G. Horney  
Office Scientific Research  
R and D Command, USAF  
Box 1395  
Baltimore, Maryland